

BZK ANTISEPTIC TOWELETTE- benzalkonium chloride swab
Guangdong Comfort Medical Products Co., Ltd.

BZK antiseptic towelette

INACTIVE INGREDIENTS

Water

DIRECTIONS

Tear open packet, unfold and use as a washcloth. Allow hands to dry without wiping.

WARNINGS

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For external use only.

DO NOT USE, in the eyes. If this happens, rinse thoroughly with water.

STOP USE and ASK A DOCTOR: If irritation or redness develop and persists for more than 72 hours.

CAUTION: Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

USES

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Antiseptic cleansing of face, hands and body to decrease bacteria on skin without soap and water.

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PURPOSE

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First Aid Antiseptic

ACTIVE INGREDIENT

Benzalkonium chloride 0.13%

Drug Facts

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OTHER INFORMATION

Store at room temperature. INACTIVE INGREDIENTS

Water.

NDC:71584-0103-1

Antiseptic Towelette

**Benzalkonium Chloride/
First Aid Antiseptic**

Dist. by:
GuangDong Comfort Medical Products Co., Ltd.
A26 District Three, Jiangmen Industrial Transfer
Industrial Park Enping District, Enping City,
Guangdong, China 529400

1 Single Use pre-
moistened pad

Drug Facts	
Active ingredients	Purpose
Benzalkonium Chloride 0.13%..	First Aid Antiseptic
Uses ■ Antiseptic cleansing of face, hands and body to decrease bacteria on skin without soap and water	
Warnings	
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Stop use and ask a doctor if irritation or redness develop and persists for more than 72 hours	
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Directions ■ tear open packet, unfold and use as a washcloth. Allow hands to dry without wiping.	
Inactive ingredients	water

Exp.Date: 04/17

Lot No.: 20140530

BZK ANTISEPTIC TOWELETTE

benzalkonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71584-0103
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71584-0103-1	1.4 mL in 1 PACKET; Type 0: Not a Combination Product	07/17/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	07/17/2017	

Labeler - Guangdong Comfort Medical Products Co., Ltd. (544507534)

Registrant - Guangdong Comfort Medical Products Co., Ltd. (544507534)

Revised: 11/2024

Guangdong Comfort Medical Products Co., Ltd.